

### **Listing of the Claims:**

The following listing of claims will replace all prior versions and listings of claims in the application.

1.-28. (Canceled)

29. (Currently Amended) A method of enhancing an antigen-specific type 1 immune response against an antigen comprising:

obtaining a pharmaceutical composition comprising an antigen, a type 1 inducing adjuvant ~~that is not an oligodeoxynucleotide (ODN) containing a CpG motif~~, and Alum; and

administering the pharmaceutical composition to a subject;

wherein an antigen-specific type 1 immune response against antigen is enhanced in the subject.

30. (Previously Presented) The method of claim 29, wherein the antigen is a viral, parasitic or bacterial antigen.

31. (Previously Presented) The method of claim 30, wherein the antigen is a viral antigen further defined as a hepatitis viral antigen, HIV-, HPV-, or influenza antigen.

32. (Previously Presented) The method of claim 31, wherein the antigen is a hepatitis viral antigen further defined as a hepatitis A, hepatitis B, hepatitis C, or hepatitis D antigen.

33. (Previously Presented) The method of claim 29, wherein the type 1 inducing adjuvant is selected from the group consisting of a polycationic polymer, lipid particle emulsions, especially MF59, stable formulations of squalene and pluronid polymers and threonyl analogs of MDP (syntex adjuvant formulation (SAF)), monophosphoryl Lipid A (MPL), saponins, especially QS21, an immunostimulatory oligodeoxynucleotide (ODN), and combinations thereof.

34. (Previously Presented) The method of claim 29, wherein the type 1 inducing adjuvant is a lipid particle emulsion further defined as MF59.

35. (Previously Presented) The method of claim 29, wherein the type 1 inducing adjuvant is a saponin further defined as QS21.

36. (Previously Presented) The method of claim 29, wherein the type 1 inducing adjuvant is an immunostimulatory ODN further defined as a deoxynucleotide comprising deoxyinosine and/or deoxyuridine residues; a deoxynucleotide comprising at least one 2'deoxycytosine-monophosphate or -monothiophosphate 3' adjacent to a 2'deoxyinosine-monophosphate or -monothiophosphate, or an ODN based on inosine and cytidine.
37. (Previously Presented) The method of claim 36, wherein the type 1 inducing adjuvant is a deoxyinosine-deoxycytosine 26-mer.
38. (Previously Presented) The method of claim 29, wherein the type 1 inducing adjuvant is a polycationic polymer further defined as a synthetic peptide containing at least 2 KKK motifs separated by a linker of 3 to 7 hydrophobic amino acids; a polycationic peptide, polylysine, or an antimicrobial peptide.
39. (Previously Presented) The method of claim 38, wherein the type 1 inducing adjuvant is a synthetic peptide with the sequence KKKLLLLLKKK.
40. (Previously Presented) The method of claim 38, wherein the type 1 inducing adjuvant is polyarginine.
41. (Previously Presented) The method of claim 38, wherein the type 1 inducing adjuvant is a cathelicidin-derived antimicrobial peptide.
42. (Previously Presented) The method of claim 29, wherein the subject is human.